

(19) World Intellectual Property Organization International Bureau





(43) International Publication Date 31 July 2003 (31.07.2003)

PCT

(10) International Publication Number WO 03/062319 A1

C08L 55/02, (51) International Patent Classification7: C08F 299/00

PCT/US03/01468 (21) International Application Number:

(22) International Filing Date: 16 January 2003 (16.01.2003)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data: 60/349,211

16 January 2002 (16.01.2002)

- · (71) Applicant (for all designated States except US): RUT-GERS, THE STATE UNIVERSITY [US/US]; Old Queens, Somerset Street, New Brunswick, NJ 08903 (US).
- (72) Inventors; and
- Inventors/Applicants (for US only): NOSKER, Thomas, J. [US/US]; 4 Green Farm Lane, Stockton, NJ 08559 (US). RENFREE, Richard, W. [US/US]; 211 Katherine Street, Scotch Plains, NJ 07076 (US).
- (74) Agents: FOLEY, Shawn, P. et al.; Lerner, David, Littenberg, Krumholz & Mentlik, LLP, 600 South Avenue, West, Westfield, NJ 07090 (US).

- (81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.
- (84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GO, GW, ML, MR, NE, SN, TD, TG).

Published:

- with international search report
- before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

10

15

20

25

30

COMPOSITIONS AND METHODS OF MAKING PLASTIC ARTICLES BACKGROUND

Plastic lumber is currently produced from high-density polyethylene ("HDPE"). This material is ubiquitous -- it is the predominant material found in non-soda bottle applications in the U.S. and can be obtained from curbside collection programs. Not surprisingly, therefore, HDPE has become an acceptable substitute for chemically treated wood in many commercial products.

If HDPE has a shortcoming in terms of performance, it is its reduced stiffness compared to wood. Standard yellow pines have a modulus (a measure of stiffness) as measured along the main axis, of 1 to 1.2 million psi. Modulus tends to decrease with duration of exposure to the environment. In the case of yellow pine, for example, over the course of 10 years of outdoor exposure, its modulus will drop by 25-50%. The modulus of HDPE-based plastic lumber is typically about 10% of new standard yellow pine. In contrast to wood however, the modulus of this type of lumber does not significantly diminish over the same period of time.

To increase the stiffness of HDPE in railroad ties, materials such as fiber-reinforced composites, polymer-fiber-reinforced composites, mineral filled polymer composites, hybrid plastics, steel and concrete composites, have been added. See U.S. Patents 5,789,477, 5,916,932 and 5,298,214. These efforts have been moderately successful, at best. It has been found to be unpredictable as to which stiffening agents can be mixed with HDPE, or any other plastic, and produce a material that is cohesive, workable and that meets the necessary performance criteria, all in a cost effective manner. For example, polyvinylchloride (PVC) and polyethylene

15

20

25

30

PCT/US03/01468

terephthalate (PET) are unworkable at conditions in which HDPE is typically processed.

In addition to identifying materials that are compatible with HDPE from these standpoints, there is recognition of the benefit of using recycled materials. This would ensure a plentiful supply of material at low cost while providing an environmentally friendly solution to waste management.

SUMMARY OF THE INVENTION

A first aspect of the present invention is directed to an immiscible polymer blend comprising (1) polyethylene (PE) and (2) acrylonitrile-butadiene-styrene (ABS), polycarbonate (PC), or a mixture of ABS and PC. The PE has a melt flow of less than about 1, and more preferably less than 1, and the PC, ABS or mixture of PC and ABS has a melt flow greater than about 1, and more preferably greater than 1. In some embodiments, the polymer blend contains PE and ABS, PE and PC, or PE, ABS and PC. In preferred embodiments, the PE is high density PE (HDPE).

A second aspect of the present invention is directed to a plastic article containing the polymer blend. In preferred embodiments, the plastic article is in the form of plastic lumber, and more preferably is in the form of a railroad tie. Methods for making the plastic articles are also provided.

PC, ABS and PC/ABS blends are typically used in the construction of the housings of electronic components and in particular consumer electronics including TVs, radios, DVD players, VCRs, CD players, stereos, video cassette cartridges, video game players and computers, including monitor housings and CPU housings, accessory components and keyboards. Many parts of a computer can be recycled. However, the housing, which accounts for a significant proportion of the size and weight of the computer, inevitably finds its way to a landfill. Due to the meteoric growth in the computer and communications industries, the number of telephones, portable

15

E.

telephones, computers and electronic devices has increased tremendously. At present, there are few recycling possibilities for these materials. Thus, a significant advantage of the present invention lies in the nature of the sources for materials needed to make it. They are plentiful, cheap and environmentally efficacious.

BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1 is a scanning electron microscopy (SEM) photograph illustrating the microstructure of an immiscible polymer blend of the present invention containing 90% PE and 10% PC, wherein the PC fibers have a high aspect ratio which is desirable for composites.

Fig. 2 is a scanning electron microscopy (SEM) photograph (at a much higher magnification) illustrating the microstructure of an immiscible polymer blend of the present invention containing 90% PE and 10% PC, wherein the diameter of the PC fibers is on the order of 30-40 nanometers, thus showing a very fine structure.

Fig. 3 is a graph that plots the modulus versus the 20 weight percent of PC in a polymer blend of the present invention containing HDPE and PC.

Fig. 4 is a graph that plots the modulus versus the weight percent of ABS in a polymer blend of the present invention containing HDPE and ABS.

25 Fig. 5 is a graph that plots the modulus versus the combined weight percent of PC and ABS in a polymer blend of the present invention containing HDPE, ABS and PC.

DETAILED DESCRIPTION OF THE INVENTION

HPPE as well as low-density polyethylene (LDPE) may be used in the polymer blends and plastic articles of the present invention. *Modern Plastics Encyclopedia* describes HDPE as having a density of more than 0.940 gm/cc, whereas LDPE is described in terms of having a density between 0.916 gm/cc and 0.940 gm/cc. Regardless of the type of PE, it has a melt flow

measured in accordance with ASTM D1238, of less than about 1 g/10 min at 190 C/2.16 Kg. In preferred embodiments, the PE has melt flow of less than 1 g/10 min at 190 C/2.16 Kg, in which embodiments the PE is said to possess a fractional melt flow. HDPE is the preferred PE for use in the present invention. HDPE can be obtained from a variety of sources; it may be virgin or recycled in nature. Recyclable HDPE is contained in commercial products such as milk bottles, water bottles and detergent bottles.

PC in accordance with the present invention 10 amongst other synthetic thermoplastic resin derived from, sources, bisphenol A and phosgene. PCs can be formed from any dihydroxy compound and carbonate di-ester or by See, e.g., U.S. Patent 5,189,139 (teaching a method of preparation of a polycarbonate comprising reacting a 15 diaryl carbonate with a bisphenol to produce a polycarbonate and a phenol, regenerating the diaryl carbonate by reacting the phenol byproduct with a carbonyl halide, and optionally recycling the hydrogen halide using a metal Redox captive 20 halogen swing reactor system). PC as used in the present invention is not limited to virgin materials; it may be recycled material. For example, PC is found in molded products, solution-cast extruded film, structural parts, tubes prosthetic devices, meter and piping, face plates, . 25 nonbreakable windows, street light globes, appliances and housings of electronic devices such as consumer electronics. Injection molding grades of PC are preferred. Methods of obtaining recycled PC are known in the art. See, The state of the search of the

ABS is an engineering plastic used in automobile body parts and for fittings in telephones, bottles, heels, luggage, packaging, refrigerator door linings, plastic pipes, building panels, shower stalls, boats, radiator grills and housings for electronics equipment and business machines including consumer

15

20

25

30

electronics. Most ABS resins are true graft polymers consisting of an elastomeric polybutadiene or rubber phase, with styrene and acrylonitrile monomers compatibility, dispersed in a rigid styrene-acrylonitrile However, mechanical poly-blends of elastomers and rigid copolymers are also available. Virgin ABS resin may be used in accordance with the present invention. somewhat expensive when supplied in its purest form. For this reason, recyclable sources of this polymer are preferred for use in the present invention. ABS is extensively used in the manufacture of inexpensive, durable products. It is a primary material in the manufacture of products such as computer housings, computer monitor housings, televisions automobile components. Methods of recycling ABS e.g., from these types of products, are known in the art. example, U.S. Patent 6,007,005. The PC, ABS and PC/ABS mixtures used in the present invention have a melt flow measured in accordance with ASTM D1238, of greater than about 1 q/10 min at 190°C/2.16 Kg. In preferred embodiments, the PC and ABS have a melt flow of greater than 1 g/10 min at 190°C/2.16 Kg.

The relative amounts of PE and the PC and/or ABS present in the polymer blends and plastic articles of the present invention vary depending on the commercial application and the desired properties. In general, the amount of PE ranges from about 50% to about 90%, and the amount of PC and/or ABS ranges from about 10% to about 50%, based upon the total weight of these polymers. As discussed below, the products of the present invention may contain additional elements. In some embodiments, the PE is present in an amount of about 60-65% by weight, and the PC and/or ABS is present in an amount of about 35-40% by weight. Beyond these general guidelines, the relative amounts of PE, PC and/or ABS will be determined on the basis of the ultimate commercial application. For

example, the strength and stiffness requirements of a formulation used for the production of decorative items, furniture, household containers and the like are not as rigorous as the requirements for railroad ties and marine pilings. Where less demanding applications are envisioned, considerations such as the availability and relative cost of the components of the blend may play a relatively important role in determining the ultimate blend composition compared to maximizing the strength and stiffness.

A significant factor in determining the relative amounts 10 of these polymers is the desired stiffness of the plastic The immiscible polymer blends of the present article. invention result in a plastic composite article having a measure of stiffness greater than that of an article made with 15 the PE alone (i.e., without the PC, ABS or mixture thereof). The stiffness of any given article must be measured in accordance with established industry standards. In the case of railroad ties, for example, the resulting modulus E of the composition should be at least about 170,000, and have strength of at least about 2500 psi, both as measured by ASTM 20 D6109-97. Referring to Fig. 3, the linear plot shows modulus of various mixtures of HDPE and ABS based on the law of The curved line shows that the blends actually mixtures. tested exhibited a modulus greater than what would have been expected based upon the additive contributions of each polymer 25 overall stiffness. Thus, a synergistic effect was The graph also shows what combinations of HDPE and achieved. ABS will result in a specific modulus, as well as which combinations will result in a stiffness greater than PE alone. 30 Here, a blend containing about 10% ABS and about 90% HDPE will have a modulus of about 175,000. Figs. 4 and 5 provide similar information and also show synergistic effects of various combinations of HDPE and ABS/PC, and HDPE and ABS, respectively. For ease of review, the data plotted on Figs.

30

3-5 are also set forth in Tables II-IV, below. Marine pilings have even higher stiffness requirements. In this case, the modulus is typically about 250,000 as measured by ASTM D6109-97.

In addition to HDPE, PC, ABS and PC/ABS blends, the blend 5 or formulations of the present invention may also include other materials, preferably in an amount of less than about 40%, most preferably less than about 20%, and even more preferably less than about 10% by weight of the article. These other materials can include other plastics or polymer 10 materials such as polyphenylene oxide (PPO). The polymer blends of the present invention preferably do not include PVC because it tends to break down during conditions under which PE is processed. Most desirably, formulations in accordance with the present invention contain less than about 1% total 15 Most preferably, the formulations useful in accordance with the present invention contain substantially no PVC (i.e., trace amounts or less). PVC is sometimes found in computer Thus, when such materials are used as sources for PC and/or ABS, the PVC should be removed. Removal of PVC can 20 be accomplished in accordance with standard techniques.

One or more reinforcing fillers may be used to further improve upon the various properties such as the tensile strength, impact strength, stiffness, and heat distortion properties of the composite. The reinforcing fillers are often used in conjunction with coupling agents, such as silanes and titanates, to facilitate the incorporation of the filler into the polymer matrix. Examples of fillers include fiberglass, asbestos, wollastonite, whiskers, carbon filaments, talc, clays (e.g., kaolin), mica, calcium carbonate, fly ash, and ceramics. Filamentous fillers such as glass fibers typically provide the greatest impact and tensile strength properties while the addition of relatively plate-like structures like talc and mica generally result in increased stiffness and heat

20

25

30

distortion. Glass fibers are desirable choices because they improve stiffness without significantly reducing impact properties or increasing density. See U.S. Patent 5,916,932. At the same time however, they are very expensive and also cause significant wear of the processing equipment. Consequently, less expensive fillers such as talc and mica have been used as substitutes for glass fibers. A potential disadvantage of these fillers, depending upon the ultimate application, is that they have a much higher density that typically results in heavier composite articles than those employing glass fibers.

The plastic articles of the present invention may contain other additives such as compatibilizers (e.g., maleic anhydride copolymers and ionomers, acrylate copolymers and ethylene acrylic acid copolymers), which effectuate the mixing (i.e., compatibility) of two or more polymers, antioxidants, UV stabilizers, lubricants, antifungal agents and colorants. These various additives may be added during fabrication of the construction article or may be present in one of the initial polymeric components. Generally, reinforcing fillers and other additives are present in an amount of less than 20% by weight of the blend or article, more preferably less than 10% by weight of the blend or article.

The immiscible polymer blends in accordance with the present invention can be used to produce a wide variety of plastic or polymer composite articles. Such articles include furniture, containers such as bottles, trash receptacles, boxes, cartons, doormats, floor coverings, lawn decorations and ornaments, automobile parts, toys, wagons and gardening carts, buildings and sheds, furniture e.g., casual furniture such as used for indoor/outdoor use including tables and chairs, park benches etc., housings for electronic components including consumer electronics such as radios, televisions, stereo systems, VCRs, DVDs, video game players and cartridges,

computer housings, keyboards and monitors and the like. Preferred plastic articles are in the form of lumber e.g., structural beams, marine pilings, railroad ties, telephone poles, blocks, boards, sheets, panels and the like. These may be structural, decorative or both. They may be textured to simulate wood grain, to simulate clapboard or cedar shake or may have raised or imprinted designs of any manner. They may Preferred articles are in the form of also include indicia. plastic lumber. Such articles may be used almost anyplace where dimensional lumber or sheets are currently used. can be used, for example, in house construction in place of boards, interior and exterior plywood, flooring, roofing, decks and rafters. More preferred applications pertain to railroad ties.

Methods of polymer processing and fabrication methods 15 known in the art for producing construction materials can be utilized to produce the blends and articles of the present These methods include both extrusion and/or invention. molding processes such as continuous extrusion, and injection and compression molding. Extrusion processes are preferred. 20 Typically, the polymer (and non-polymeric) components are dry blended at about room temperature, and then fed into an extruder where they are melted and mixed. The operating conditions of the extruder may vary but an L (length) to D (diameter) ratio of from about 24:1 to about 25 compression screw ratio of about 3:1, and an extrusion temperature of about 220 °C are preferred. The extruded blend comes out of a die into a mold, shaped in the form of the article (or a part thereof). In the case of a railroad tie, for example, the mold typically has dimensions of about 7" x 30 $9" \times 8.5-10'$. Once the filled mold is cooled to about room temperature, its contents are ejected.

Further steps may be necessary or desirable in certain circumstances, such as when the blend contains PC or when the

composition of the plastics (e.g., a recycled source) unknown. If the blend includes PC, which is hydroscopic in nature, it is important that it be dried at some time before being melted. Dessication dryers, vacuum ovens and the like are suitable for this purpose. If the content of the incoming plastic material is not known, then a sorting step should be One method of sorting in accordance with the employed. present invention utilizes a Fourier transform infrared spectrophotometer ("FTIR") set in reflective mode. The FTIR may have a photo acoustic detector attachment. A database is 10 prepared of the FTIR patterns of as many polymer types as possible. Then the patterns of unknown materials can be matched to the database of known samples. Samples of unknown materials can be obtained by producing a small disc punched from their bodies. Those punched sections can be analyzed by 15 FTIR. Most preferably, the analysis is conducted by exposing the cross section of the disc to the FTIR such that the results are not distorted in the event that the inner or outer layer of the source is covered by a material different than 20 the core. In this way the composition of the incoming raw material can be analyzed. X-ray fluorescence spectrophotometry is another useful technique. This technique is advantageously employed to detect PVC. The object of such sorting is two-fold, namely: 1) to identify the materials 25 going into the polymer blends so as to predict the resulting properties; and 2) to eliminate undesirable components such as This sorting technique is principally used to identify PVC. the composition of PC, ABS or PC/ABS blends in an incoming Note: The state of 30 ... used in connection with determining whether unwanted materials such as PVC are present and/or in connection with the PE.

Depending on the content of PC, ABS or PC/ABS blend contained in a recyclable waste material, it may be necessary or desirable to augment with additional quantities of one or

more of the plastic materials. After sorting, the resulting materials can be stored for later use or processed and formed as discussed above.

Class 1 railroad ties are a particularly demanding 5 application. Railroad ties are exposed to significant stress. Not only must they bear the weight of a train but they must also bear the dynamic stress of a moving train. They must withstand extremes of temperature and moisture as well as environmental factors such as oil, diesel fuel, acid rain and the like. See Nosker et al., Development of a Recycled 10 Plastic/Composite Cross Tie, 98 Am. Railway Engineering Bulletin No. 760 (1997) and Nosker et al., A Performance-Based Approach to the Development of Recycled Plastic Composite Cross Tie, Proc. Society of Plastic Engineers 1998 Antec Conference, Atlanta, GA, April 1998. Properties of railroad 15 ties (e.g., testing requirements) and methods of making railroad ties from plastics, including recycled plastic materials, are known in the art. See, U.S. Patents 5,569,424; 5,789,477; 5,886,078; 5,916,932; and 6,191,228 (teaching the "batch" and "continuous" processes). 20

The invention will now be described by way of the following examples. These examples are not meant to limit the invention in any way. Unless otherwise indicated, all parts and percentages are by weight. Tables I-IV below compare the tensile properties of pure, bottle grade, recycled HDPE, ABS, PC and mixtures of PC/ABS with various blends, as measured in accordance with ASTM D638.

TABLE I - Tensile properties of HDPE

SAMPLE	MODULUS (psi)	ULTIMATE STRENGTH (psi)	
100% HDPE	118,000	2,600	

TABLE II - Tensile properties of mixtures of ABS and HDPE

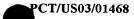
SAMPLE	MODULUS (psi)	ULTIMATE STRENGTH (psi)	
10% ABS 90% HDPE	128,000	2,700	
20% ABS 80% HDPE	145,000	2,800	
30% ABS 70% HDPE	218,000	3.300	
35% ABS 65% HDPE	229,000	3,400	
40% ABS 60% HDPE	222.000	3,500	
100% ABS	390,000	5,500	

TABLE III - Tensile properties of mixtures of PC/ABS and HDPE

SAMPLE	MODULUS (psi)	ULTIMATE STRENGTH (psi)		
10% PC/ABS 90% HDPE	186,000 3,100			
20% PC/ABS 80% HDPE	207,000	3,300		
30% PC/ABS 70% HDPE	228,000	3,700		
35% PC/ABS 65% HDPE	250,000	4,100		
40% PC/ABS 60% HDPE	253,000	4,300		
100% PC/ABS	418,000	7,900		
VACUUMED PC/ABS	419,000	8,000		

TABLE IV - Tensile properties of mixtures of PC and HDPE

SAMPLE	SAMPLE MODULUS ULTIM (psi)			
10% PC 90% HDPE	179,000	3,100		
20% PC 80% HDPE	226,000	3,400		



30% PC 70% HDPE	213,000	3,800
35% PC 65% HDPE	234,000	4.100
40% PC 60% HDPE	274,000	4,200
PC DRY	408,000	9,000

Tables V-VII compare the modulus and ultimate strength of recycled materials versus their virgin counterparts. The data show that the properties are comparable such that virgin or recycled polymers can be used.

TABLE V - Tensile properties of 100% recycled ABS and virgin ABS

SAMPLE	MODULUS (psi)	ULTIMATE STRENGTH (psi) 5,500 3,300-8,000	
100% ABS	390,000		
Virgin ABS	270,000-400,000		

TABLE VI - Tensile properties of 100% recycled PC/ABS and 10 virgin PC/ABS

SAMPLE	MODULUS ULTIMATE STR (psi) (psi)	
100% PC/ABS	418,000	7,900
VACUUMED 100% PC/ABS	C/ABS 419,000 8,000	
Virgin PC/ABS	350,000-455,000	5,800-9,3000

TABLE VII - Tensile properties of 100% recycled PC and virgin PC

SAMPLE	MODULUS (psi)	ULTIMATE STRENGTH (psi)	
100% DRY PC	408,000	9,000	
Virgin PC	350,000	9,200-9,500	

15

All patent and non-patent publications cited in this specification are indicative of the level of skill of those skilled in the art to which this invention pertains. All these publications and patent applications are herein incorporated by reference to the same extent as if each individual publication or patent application was specifically and individually indicated as being incorporated by reference herein.

Although the invention herein has been described with reference to particular embodiments, it is to be understood that these embodiments are merely illustrative of the principles and applications of the present invention. It is therefore to be understood that numerous modifications may be made to the illustrative embodiments and that other arrangements may be devised without departing from the spirit and scope of the present invention as defined by the appended claims.

CLAIMS:

- 1. An immiscible polymer blend comprising polyethylene (PE) and acrylonitrile-butadiene-styrene (ABS) or polycarbonate (PC) or a mixture of ABS and PC, wherein said PE has a melt flow at 190°C/2.16Kg of less than about 1, and said PC, ABS or mixture of PC and ABS has a melt flow at 190°C/2.16Kg greater than about 1.
- 2. The polymer blend of claim 1 which comprises PE and ABS.
- The polymer blend of claim 1 which comprises PE and PC.
 - 4. The polymer blend of claim 1 which comprises PE, ABS and PC.
- 5. The polymer blend of claim 1 wherein said PE is a 15 high density PE (HDPE).
 - 6. The polymer blend of claim 1 wherein said PE is a low density PE (LDPE).
 - 7. The polymer blend of claim 1 wherein said PE has a fractional melt flow.
- 20 8. The polymer blend of claim 1 wherein said PC, ABS or mixture of PC and ABS has a melt flow greater than 1.
 - 9. The polymer blend of claim 1 wherein said PE has a fractional melt flow and wherein said PC, ABS or mixture of PC and ABS has a melt flow greater than 1.
- 25 10. The polymer blend of claim 9 wherein said PE is HDPE.
 - 11. A plastic or polymer composite article formed of the polymer blend of claim 1.
 - 12. The plastic article of claim 11 which is lumber.
- 30 13. The plastic article of claim 11 which is a railroad tie.
 - 14. The plastic article of claim 11 which is a marine piling.



- 15. A method of making a plastic or polymer composite article, comprising:
- (a) preparing an immiscible polymer blend comprising polyethylene (PE) and acrylonitrile-butadiene-styrene (ABS) or polycarbonate (PC) or a mixture of ABS and PC, wherein said PE has a melt flow at 190°C/2.16Kg of less than about 1, and said PC, ABS or mixture of PC and ABS has a melt flow at 190°C/2.16Kg greater than about 1; and
- (b) shaping the blend into a desired shape of the 10 article.
 - 16. The method of claim 15 wherein said preparing and shaping comprise continuous extrusion.
 - 17. The method of claim 15 wherein said preparing comprises extrusion.
- 15 18. The method of claim 15 wherein said shaping comprises molding.
 - 19. The method of claim 15 wherein said preparing and shaping comprises injection molding.



PC/PE 10/90 Perpendicular

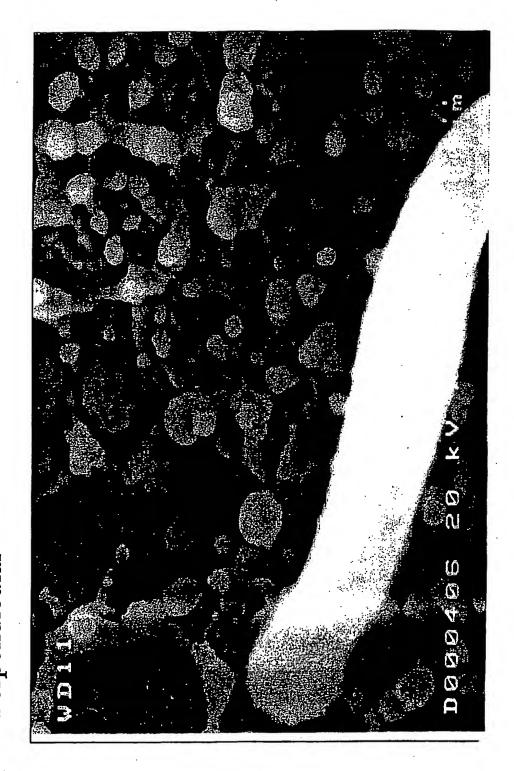


Fig. 2

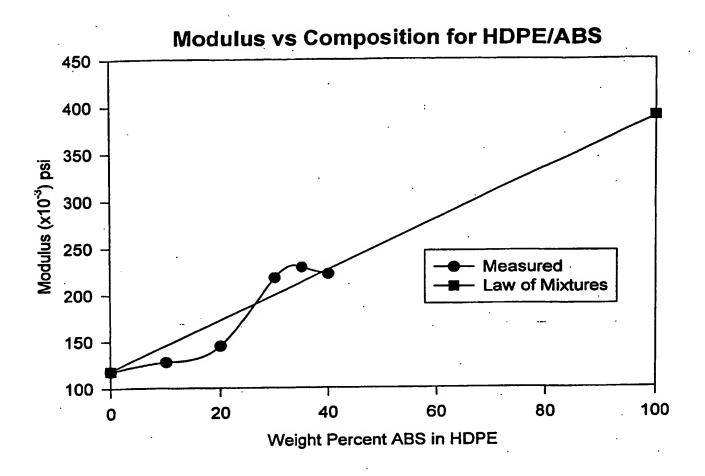


Fig. 3

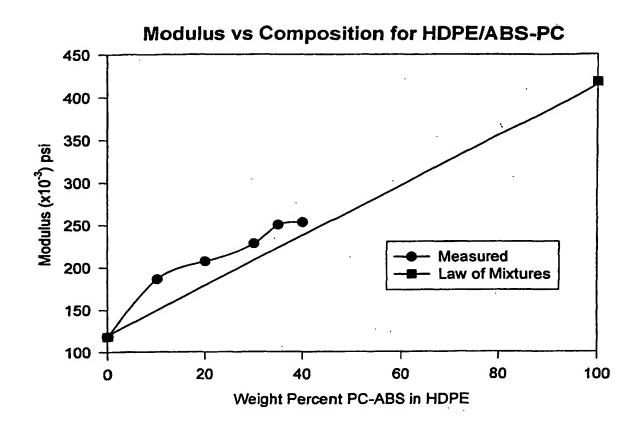


Fig. 4

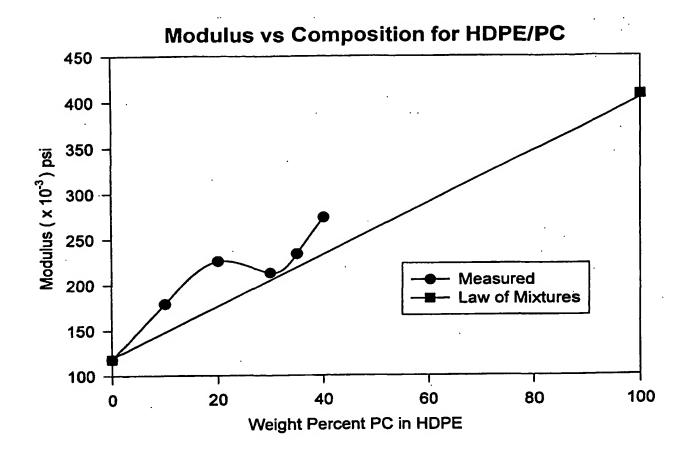


Fig. 5



INTERNATIONAL SEARCH REPORT

International application No.

PCT/US03/01468

	A. CLASSIFICATION OF SUBJECT MATTER IPC(7) : C08L 55/02, C08F 299/00					
	US CL : 525/70, 185					
	According to International Patent Classification (IPC) or to both national classification and IPC B. FIELDS SEARCHED					
	Minimum documentation searched (classification system followed by classification symbols) U.S.: 525/70, 185					
	Documentation	Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched				
				-		
	Electronic da	ta base consulted during the international search (name	of data t	pase and, where practicable, sear	rch terms used)	
	C. DOC	UMENTS CONSIDERED TO BE RELEVANT				
	Category *	Citation of document, with indication, where a			Relevant to claim No.	
	X 	US 4,110,420 A (TURNER) 29 August 1978 (29.08	.1978), A	bstract.	1, 2, 5-11, 14-19	
	A				3, 4, 12, 13	
	x 	US 5,098,610 A (OOAMURA et al.) 24 March 1993 lines 63-66.	2 (24.03.	1992), Abstract, column 3,	1, 3-11	
	A	mes 03-00.			2, 12-19	
	,					
	Further	documents are listed in the continuation of Box C.		See patent family annex.		
•	· s	pecial categories of cited documents:	-T-	later document published after the inte date and not in conflict with the appli		
		defining the general state of the art which is not considered to be that relevance	-x-	principle or theory underlying the inv document of particular relevance; the		
		plication or patent published on or after the international filing date	^	considered novel or cannot be considered when the document is taken alone		
		which may throw doubts on priority claim(s) or which is cited to the publication date of another citation or other special reason (as	-Y"	document of particular relevance; the considered to involve an inventive ste		
		referring to an oral disclosure, use, exhibition or other means		combined with one or more other suc being obvious to a person skilled in the	h documents, such combination	
programme to the section of the sect		published prior to the international filing date but later than the ate claimed	-&"	document member of the same patent	family	
	٠. ٠	ctual completion of the international search		mailing of the international sear MAY 2003	rch report	
		3 (30.04.2003) ailing address of the ISA/US	l	zed officer		
	Ma	I Stop PCT, Attn: ISA/US	·	Jeffrey .	0.2	
grander vielenden der	P.O	nmissioner for Patents . Box 1450			7	
		xandria, Virginia 22313-1450 . (703)305-3230	reiepno	one No. 703 306 5665	<i>J</i> .	
		N/210 (second sheet) (July 1998)	·		····	
i er vileni e						
		•				
· · · · · · · · · · · · · · · · · · ·						
and the second of the second o	a garanda kanada Pagana da					